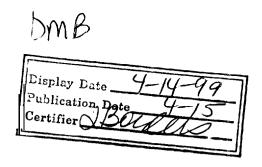
DEPARTMENT OF HEALTH AND HUMAN SERVICES



Food and Drug Administration

21 CFR Parts 201,330,331,341,346, 355,358,369, and 701

[Docket Nos. 98 N-0337, 96 N-0420, 95 N-0259, and 90P-0201]

RIN 0910-AA79

Over-The-Counter Human Drugs; Labeling Requirements; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of March 17, 1999 (64 FR 13254). The final rule established a standardized format and standardized content requirements for the labeling of over-the-counter (OTC) drug products. The document was inadvertently published with an incorrect effective date. This document corrects that error.

DATES: Effective (*insert date of publication*). The effective date of the final rule published on March 17, 1999 (64 FR 13254), is corrected to be May 16, 1999.

FOR FURTHER INFORMATION CONTACT: Debra L. Bowen, Food and Drug Administration, Center for Drug Evaluation and Research (HFD–560), 5600 Fishers Lane, Rockville, MD 20852, 301–827–2222, or email ''BOWEND@cder.fda. gov".

SUPPLEMENTARY INFORMATION: In FR Dec. 99–6296, appearing on page 13254 in the **Federal Register** of Wednesday, March 17, 1999, the following correction is made:

1. On page 13254, in the first column, under the "Dates" section 'Effective Date: April 16, 1999." is corrected to read "Effective Date: May 16, 1999."

Dated: 4/12/99 April 12, 1999

William K. Hubbard

Acting Deputy Commissioner for Policy

William Hustand

[FR Dec. 99-???? Filed ??-??-99; 8:45 am]

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